## Claims:

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- 1. A method of inducing an immune response by the delivering of an effective amount of lipid-tailed protein to a mucosal membrane of a subject.
- 2. The method of Claim 1, wherein the lipoprotein is applied to the mucosal membrane without adjuvant.
- 3. The method of Claim 1, wherein the lipoprotein is applied to the mucosal membrane without using a needle.
- 4. The method of Claim 1, wherein the lipoprotein is applied intranasally, sublingually, by eye-drops, or suppositories.
- 5. The method of Claim 1, wherein the lipoprotein has at least one lipid coupled to a functional group of the said protein.
- 6. The method of Claim 1, wherein the lipoprotein has at least one lipid coupled to a  $\alpha$ -NH<sub>2</sub> and/or  $\epsilon$ -NH<sub>2</sub> functional group of the peptide.
- 7. The method of Claim 1, wherein application of the lipoprotein induces a B cell response.
- 8. The method of Claim 1, wherein application of the lipoprotein induces a T cell response.
- 9. The method of Claim 1, wherein application of the lipoprotein induces a systemic B and/or T cell response.
- 10. A composition consisting in at least one lipoprotein inducing a mucosal immune response in vivo in absence of toxic adjuvant.

- 11. A composition according to Claim 10, wherein the adjuvant is non-toxic for the mucosal membranes.
  - 12. A lipopeptide, wherein the lipopeptide is tailed with a lipid component.
- 13. The lipopeptide of Claim 11, wherein the lipid component is a palmitoyl residue having 16 carbon atoms.
  - 14. The lipopeptide of Claim 12, wherein the lipopeptide is: LSA3-NRII Ac-LEESQVNDDIFNSLVKSVQQEQQHNVK(PAM)NH2 OR LSA1-J Ac-ERRAKEKLQEQQSDLEQRKADTKKK(PAM).
- 15. The method of Claim 9, wherein the lipopetide is:
  LSA3-NRII Ac-LEESQVNDDIFNSLVKSVQQEQQHNVK(PAM)NH2 OR
  LSA1-J Ac-ERRAKEKLQEQQSDLEQRKADTKKK(PAM)NH2.

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- 16. A composition consisting in at least one lipopeptide inducing a mucosal immune response *in vivo* in the absence of toxic adjuvant, wherein the lipopeptide is at least one lipopeptide according to Claim 13.
- 17. A vaccine composition for mucosal administration containing at least one lipopeptide inducing an B and/or T cell response in vivo in absence of adjuvant.
- 18. A vaccine composition containing a lipopeptide according to Claim 13 in the absence of adjuvant.
  - 19. An immunogenic composition containing a lipopeptide according to Claim 13.
- 20. A method of stimulating T-Lymphocyte responses in vitro after immunization via mucosal administration comprising the following steps:
  - a) immunizing BALB/C mice by mucosal administration with a peptide tetanic toxinpol HIV palmitic antigen,

- b) collecting of ganglia sub-mandibulaires at day 15, and
- c) visualizing T cell responses by labeling target cells with CFSE.
- 21. The method of Claim 1, further comprising administering a composition containing a lipid-tailed polypeptide or peptide, said lipid-tailed peptide having at least a lipid residue bound to an epitope T amino acid sequence and optionally at least one epitope B amino acid sequence.
- 22. The method of Claim 21, wherein the lipopeptide is an antigenic lipopeptide of sequence:

H-K(PAM)TT-pol 476-484

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- Nh2-K(N∈Pam)GRQYIKKANSKFIGITERGRILKEP-COOH.
  - 23. The method of Claim 1, wherein the lipopeptide is a lipid-tailed epitope T.
- 24. The method of Claim 23, wherein the lipopeptide is a lipid-tailed epitope T covalently linked to an epitope B peptide.
- 25. A composition comprising lipid-tailed polypeptide or peptide, said lipid-tailed peptide having at least a lipid residue bound to an epitope T amino acid sequence and optionally at least one epitope B amino acid sequence.

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